CLAIMS

1. A compound according to formula I

$$(R^{1})_{m}$$
 P
 X^{2}
 X^{1}
 X^{7}
 X^{6}
 $(R^{3})_{p}$
 $X^{3-}X^{4}$
 Q
 $(R^{2})_{n}$
 (I)

5 wherein

10

15

20

P is selected from aryl and heteroaryl;

 R^1 is attached to P via a carbon atom on ring P and is selected from the group consisting of hydroxy, halo, nitro, $C_{1\text{-}6}$ alkylhalo, $OC_{1\text{-}6}$ alkylhalo, $C_{1\text{-}6}$ alkyl, $OC_{1\text{-}6}$ alkyl, $C_{2\text{-}6}$ alkenyl, $C_{2\text{-}6}$ alkynyl, $OC_{2\text{-}6}$ alkynyl, $C_{0\text{-}6}$ alkyl $C_{3\text{-}6}$ cycloalkyl, $C_{0\text{-}6}$ alkylaryl, $OC_{0\text{-}6}$ alkylaryl, $C_{0\text{-}6}$ alkyl $C_{3\text{-}6}$ cycloalkyl, $C_{0\text{-}6}$ alkylaryl, $OC_{0\text{-}6}$ alkylaryl, CHO, $(CO)R^5$, $O(CO)R^5$, $O(CO)OR^5$, $O(CNR^5)OR^5$, $C_{1\text{-}6}$ alkyl CO_2R^5 , $C_{2\text{-}6}$ alkyl CO_2R^5 , $C_{1\text{-}6}$ alkyl CO_2R^5 , $C_{1\text{-}6}$ alkyl CO_2R^5 , $C_{0\text{-}6}$ alkyl CO_2R^5 , $C_{0\text{-}6}$ alkyl CO_3R^5 , $C_{0\text{-}6}$ alkyl

X¹ is selected from the group consisting of: N, NR⁴ and CR⁴;

X² is selected from the group consisting of: C and N;

X³ is selected from the group consisting of: CR⁴, N and O;

X⁴ is selected from the group consisting of: CR⁴, N, NR⁴ and O;

 X^5 is selected from the group consisting of: a bond, CR^4R^4 ', NR^4 , O, S, SO and SO₂; X^6 is selected from the group consisting of: CR^4 and N;

X⁷ is selected from the group consisting of: C and N;

5

10

15

 R^4 is independently selected from a group consisting of hydrogen, hydroxy, $C_{1\text{-}6}$ alkyl, $C_{0\text{-}6}$ alkylcyano, oxo, =NR⁵, =NOR⁵, $C_{1\text{-}4}$ alkylhalo, halo, $C_{3\text{-}7}$ cycloalkyl, $O(CO)C_{1\text{-}4}$ alkyl, $C_{1\text{-}4}$ alkyl(SO) $C_{0\text{-}4}$ alkyl, $C_{1\text{-}4}$ alkyl(SO) $C_{0\text{-}4}$ alkyl, $C_{1\text{-}4}$ alkyl, $C_{1\text{-}4}$ alkylNR⁵ and $C_{0\text{-}4}$ alkylNR⁵R⁶;

Q is selected the group consisting of heterocycloalkyl and heteroaryl;

R² and R³ are independently selected from the group consisting of: hydrox y, C₀.

6alkylcyano, oxo, =NR⁵, =NOR⁵, C₁₋₄alkylhalo, halo, C₁₋₆alkyl, C₃₋₆cycloal kyl, C₀.

6alkylaryl, C₀₋₆alkylheteroaryl, C₁₋₆alkylcycloalkyl, C₀₋₆alkylheterocycloalkyl, OC₁₋₄alkyl,

OC₀₋₆alkylaryl, O(CO)C₁₋₄alkyl, (CO)OC₁₋₄alkyl, C₀₋₄alkyl(S)C₀₋₄alkyl, C₁₋₄alkyl(SO)C₀.

4alkyl, C₁₋₄alkyl(SO₂)C₀₋₄alkyl, (SO)C₀₋₄alkyl, (SO₂)C₀₋₄alkyl, C₁₋₄alkylOR⁻⁵, C₀.

4alkylNR⁵R⁶ and a 5- or 6-membered ring containing atoms independently selected from C,

N, O and S, which ring may optionally be fused with a 5- or 6-membered ring containing atoms independently selected from the group consisting of C, N and O and wherein said ring and said fused ring may be substituted by one or more A;

wherein any $C_{1\text{-}6}$ alkyl, aryl, or heteroaryl defined under R^1 , R^2 and R^3 may be substituted by one or more A;

A is selected from the group consisting of: hydrogen, hydroxy, halo, nitro, oxo, C₀.

6alkylcyano, C₀₋₄alkylC₃₋₆cycloalkyl, C₁₋₆alkyl, -OC₁₋₆alkyl, C₁₋₆alkylhalo, OC₁₋₆alkylhalo,
C₂₋₆alkenyl, C₀₋₃alkylaryl, C₀₋₆alkylOR⁵, OC₂₋₆alkylOR⁵, C₀₋₆alkylSR⁵, OC₂₋₆alkylSR⁵,
(CO)R⁵, O(CO)R⁵, OC₂₋₆alkylcyano, OC₁₋₆alkylCO₂R⁵, O(CO)OR⁵, OC₁₋₆alkyl(CO)R⁵, C₁₋₆alkyl(CO)R⁵, NR⁵OR⁶, C₀₋₆NR⁵R⁶, OC₂₋₆alkylNR⁵R⁶, C₀₋₆alkylNR⁵R⁶, OC₁₋₆alkylNR⁵R⁶, OC₂₋₆alkylNR⁵(CO)R⁶, C₀₋₆alkylNR⁵(CO)NR⁵R⁶,
O(CO)NR⁵R⁶, OC₂₋₆alkylNR⁵(CO)R⁶, C₀₋₆alkylNR⁵(SO₂)R⁶, OC₂₋₆alkylNR⁵(SO₂)R⁶, OC₂₋₆alkylNR⁵(SO₂

₆alkyl(SO₂)R⁵, C₀₋₆alkyl(SO)R⁵, OC₂₋₆alkyl(SO)R⁵ and a 5- or 6-membered ring containing atoms independently selected from the group consisting of C, N, O and S;

R⁵ and R⁶ are independently selected from, H, C₁₋₆alkyl, C₃₋₇cycloalkyl and aryl;

m is selected from 0, 1, 2, 3 or 4;

n is selected from 0, 1, 2, 3 or 4;

p is selected from 0, 1, 2, 3 or 4; and

a salt or hydrate thereof,

with the proviso that the compound is not:

- 4,4'-(1,2-piperazinediyl)di-antipyrine;
- 4,4'-(1,2-piperazinediyl)di-antipyrine dihydrochloride; or
 - 4,4'-(1,2-piperazinediyl)di-antipyrine dipicrate;
 - 2. A compound according to claim 1 wherein m is selected from 1, 2, 3 or 4
 - 3. A compound according to claim 1 wherein X^7 is C.
- 4. A compound according to claim 1 wherein X⁵ is selected from the group consisting of CR⁴R⁴, NR⁴, O, S, SO and SO₂.
 - 5. A comound according to claim 1 wherein X^3 is selected from the group consisting of N and O.
 - 6. A compound according to claim 1 wherein P is aryl.
 - 7. A compound according to claim 6 wherein P is phenyl.
- 8. A compound according to claim 7 wherein m is selected from the group constisting of 1 and 2.
 - 9. A compound according to claim 1 wherein R^1 is selected from the group consisting of: halo, C_{1-6} alkylhalo, OC_{1-6} alkylhalo, OC_{1-6} alkylhalo, OC_{1-6} alkylhalo, OC_{1-6} alkyl $OC_{$

10. A compound according to claim 9 wherein R¹ is selected from the group consisting of: Cl, F, Me, OMe, CF₃, OCF₃, and CN.

- 11. A compound according to claim 1 wherein X^2 is C.
- 12. A compound according to claim 11 wherein X¹ is N or CR⁴.
- 13. A compound according to claim 12 wherein when X^3 is O, X^4 is N and when X^3 is N, X^4 is O.
 - 14. A compound according to claim 1 wherein X^2 is N.
 - 15. A compound according to claim 14 wherein X¹ is N.
 - 16. A compound according to claim 15 wherein X³ is N and X⁴ is N or CR⁴.
- 10 17. A compound according to claim 1 wherein X^6 is N.
 - 18. A compound according to claim 12 wherein X^5 is selected from the group consisting of a bond, $CR^4R^{4'}$, NR^4 and O.
 - 19. A compound according to claim 13 wherein X⁵ is selected from the group consisting of a bond, O and NR⁴.
- 20. A compound according to claim 16 wherein X⁵ is selected from the group consisting of O and CR⁴.
 - 21. A compound according to claim 1 wherein R^4 is selected from the group consisting of: hydrogen, C_{1-6} alkyl, C_{1-6} alkylhalo and halo.
 - 22. A compound according to claim 1 wherein Q is heteroaryl.
- 23. A compound according to claim 1 wherein Q is selected from the group consisting of:

24. A compound according to claim 23 wherein Q is

10

25. A compound according to claim 1 wherein R^2 and R^3 are independently selected from the group consisting of: C_{1-4} alkylhalo, C_{1-6} alkyl, C_{3-6} cycloalkyl, C_{0-6} alkylaryl and C_{0-6} alkylheteroaryl.

- 5 26. A compound according to claim 1 wherein A is selected from the group consisting of: hydrogen, hydroxyl, halo, C₀₋₆alkylcyano, C₁₋₆alkyl, -OC₁₋₆alkyl, C₁₋₆alkylhalo, OC₁₋₆alkylhalo.
 - 27. A compound according to claim 1 selected from:

4-(5-{2-[5-(3-Chloro-phenyl)-isoxazol-3-yl]-piperidin-1-yl}-4-methyl-4H [1,2,4]triazol-3-yl)-pyridine

3-[5-(3-Chloro-phenyl)-isoxazol-3-yl]-4-(4-methyl-5-pyridin-4-yl-4H-[1,2,4]triazol-3-yl)-morpholine

3-[5-(3-Chloro-phenyl)-isoxazol-3-yl]-4-[5-(4-difluoromethoxy-phenyl)-4-methyl-4H-[1,2,4]triazol-3-yl]-morpholine

3-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-4-(4-methyl-5-pyridin-4-yl-4H-[1,2,4]triazol-3-yl)-morpholine

3-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-4-[5-(4-difluoromethoxy-phenyl)-4-methyl-4H-[1,2,4]triazol-3-yl]-morpholine

3-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-4-(4-methyl-5-pyridin-4-yl-4H-

20 [1,2,4]triazol-3-yl)-piperazine-1-carboxylic acid tert-butyl ester

2-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-1-(4-methyl-5-pyridin-4-yl-4H-1,2,4]triazol-3-yl)-piperazine

2-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-4-methyl-1-(4-methyl-5-pyridin-4-yl-4H-[1,2,4]triazol-3-yl)-piperazine

3-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-4-[5-(4-difluoromethoxy-phenyl)-4-methyl-4H-[1,2,4]triazol-3-yl]-piperazine-1-carboxylic acid tert-butyl ester

- 2-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-1-[5-(4-difluoromethoxy-phenyl)-4-methyl-4H-[1,2,4]triazol-3-yl]-piperazine
- 5 2-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-1-[5-(4-difluoromethoxy-phenyl)-4-methyl-4H-[1,2,4]triazol-3-yl]-4-methyl-piperazine
 - 2-[2-(3-Chlorophenyl)-2H-tetrazol-5-yl]-1-{5-[4-(difluoromethoxy)phenyl]-4-methyl-4H-1,2,4-triazol-3-yl}piperidine
- 4-(5-{2-[2-(3-chlorophenyl)-2H-tetrazol-5-yl]piperidin-1-yl}-4-methyl-4H-1,2,4-triazol-3-yl)pyridine
 - 2-[2-(3-Chlorophenyl)-2H-tetrazol-5-yl]-1-[5-(4-methoxyphenyl)-4-methyl-4H-1,2,4-tria-zol-3-yl]piperidine
 - $[4-(5-\{2-[2-(3-chlorophenyl)-2H-tetrazol-5-yl]piperidin-1-yl\}-4-methyl-4H-1,2,4-triazol-3-yl)phenyl] dimethylamine$
- [4-(5-{2-[2-(3-Chloro-phenyl)-2H-tetrazol-5-yl]-piperidin-1-yl}-4-methyl-4H-[1,2,4]triazol-3-yl)-benzyl]-dimethyl-amine

20

- {2-[4-(5-{2-[2-(3-Chloro-phenyl)-2H-tetrazol-5-yl]-piperidin-1-yl}-4-methyl-4H-[1,2,4]triazol-3-yl)-phenoxy]-ethyl}-dimethyl-amine
- (R)-3-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-4-(4-methyl-5-pyridin-4-yl-4H-[1,2,4]triazol-3-yl)-morpholine
- (S) 3-[3-(3-Chloro-phenyl)-[1,2,4]oxadiazol-5-yl]-4-(4-methyl-5-pyridin-4-yl-4H-[1,2,4]triazol-3-yl)-morpholine
 - $\label{eq:continuous} $$(R)-2-[2-(3-Chlorophenyl)-2H-tetrazol-5-yl]-1-\{5-[4-(difluoromethoxy)phenyl]-4-methyl-4H-1,2,4-triazol-3-yl\} piperidine$
- 25 (S)-2-[2-(3-Chlorophenyl)-2H-tetrazol-5-yl]-1-{5-[4-(difluoromethoxy)phenyl]-4-methyl-4H-1,2,4-triazol-3-yl}piperidine

(R)-4-(5-{2-[2-(3-Chlorophenyl)-2H-tetrazol-5-yl]piperidin-1-yl}-4-methyl-4H-1,2,4-tria-zol-3-yl)pyridine

- $(S)-4-(5-\{2-[2-(3-Chlorophenyl)-2H-tetrazol-5-yl]piperidin-1-yl\}-4-methyl-4H-1,2,4-triazol-3-yl)pyridine \\$
- 4-[5-(5-{2-[5-(3-Chloro-phenyl)-isoxazol-3-yl]-pyrrolidin-1-yl}-4-cyclopropyl-4H-[1,2,4]triazol-3-yl)-pyridin-2-yl]-morpholine,
 - 4-[5-(5-{2-[5-(3-Chloro-phenyl)-isoxazol-3-yl]-pyrrolidin-1-yl}-4-methyl-4H-[1,2,4]triazol-3-yl)-pyridin-2-yl]-morpholine,
- 3-(5-{2-[5-(3-Chloro-phenyl)-isoxazol-3-yl]-pyrrolidin-1-yl}-4-methyl-4H-[1,2,4]triazol-3-yl)-pyridine,
 - 4-(5-{2-[5-(3-Chloro-phenyl)-isoxazol-3-yl]-pyrrolidin-1-yl}-4-cyclopropyl-4H-[1,2,4]triazol-3-yl)-pyridine,
 - 3-[5-(3-Chloro-phenyl)-[1,2,4]oxadioazol-3-yl]-4-(5-pyridin-4-yl-4H-[1,2,4]triazol-3-yl)-morpholine,
- 3-[5-(3-chlorophenyl)isoxazol-3-yl]-4-(4- cyclopropyl-5-pyridin-3-yl-4H-1,2,4-triazol-3-yl)morpholine,
 - 3-[5-(3-chlorophenyl)isoxazol-3-yl]-4-(4- cyclopropyl -5-pyridin-4-yl-4H-1,2,4-triazol-3-yl)morpholine,
- 3-[5-(3-chlorophenyl)isoxazol-3-yl]-4-(4-methyl-5-pyridin-3-yl-4H-1,2,4-triazol-3-yl)morpholine,
 - 3-[5-(3-Chloro-phenyl)-isoxazol-3-yl]-4-[5-(6-methoxy-pyridin-3-yl)-4-methyl-4H-[1,2,4]triazol-3-yl]-morpholine,
 - 3-[3-(3-chlorophenyl)-1,2,4-oxadiazol-5-yl]-4-[5-(2-methoxypyridin-4-yl)-4-methyl-4H-1,2,4-triazol-3-yl]morpholine,
- 3-[3-(3-chlorophenyl)-1,2,4-oxadiazol-5-yl]-4-[5-(2-methylpyridin-4-yl)-4-methyl-4H-1,2,4-triazol-3-yl]morpholine,

- 3-[3-(3-chlorophenyl)-1,2,4-oxadiazol-5-yl]-4-[5-(5-fluoropyridin-3-yl)-4-methyl-4H-1,2,4-triazol-3-yl] morpholine,
- 3-[5-(3-chlorophenyl)isoxazol-3-yl]-4-[5-(5-fluoropyridin-3-yl)-4-methyl-4H-1,2,4-triazol-3-yl]morpholine,
- ⁵ 3-[3-(3-chlorophenyl)-1,2,4-oxadiazol-5-yl]-4-(4-methyl-5-pyridin-2-yl-4H-1,2,4-triazol-3-yl)morpholine,
 - 4-[5-(5-fluoropyridin-3-yl)-4-methyl-4H-1,2,4-triazol-3-yl]-3-[3-(3-iodophenyl)-1,2,4-oxadiazol-5-yl]morpholine,
- 3-[3-(3-iodophenyl)-1,2,4-oxadiazol-5-yl]-4-(4-methyl-5-pyridin-4-yl-4H-1,2,4-triazol-3-yl)morpholine,
 - 3-[5-(3-chlorophenyl)isoxazol-3-yl]-4-[5-(2-methylpyridin-4-yl)-4-methyl-4H-1,2,4-triazol-3-yl]morpholine,
 - 3-[2-(3-chlorophenyl)-2H-tetrazol-5-yl]-4-(4-methyl-5-pyridin-3-yl-4H-1,2,4-triazol-3-yl)morpholine,
- 3-[2-(3-chlorophenyl)-2H-tetrazol-5-yl]-4-[5-(3,5-difluorophenyl)-4-methyl-4H-1,2,4-triazol-3-yl]morpholine,
 - $3-(5-\{2-[5-(3-chlorophenyl)isoxazol-3-yl]pyrrolidin-1-yl\}-4-cyclopropyl-4H-1,2,4-triazol-3-yl)pyridine, and$
 - 4-(5-{2-[5-(3-chlorophenyl)isoxazol-3-yl]pyrrolidin-1-yl}-4-methyl-4H-1,2,4-triazol-3-yl)pyridine.

20

25

- 28. A pharmaceutical composition comprising as active ingredient a therapeutically effective amount of the compound according to any one of claims 1 to 26, in association with one or more pharmaceutically acceptable diluent, excipients and/or inert carrier.
- 29. The pharmaceutical composition according to claim 28, for use in the treatment of mGluR 5 mediated disorders.
 - 30. The compound according to any one of claims 1 to 27, for use in therapy.

31. The compound according to any one of claims 1 to 27, for use in treatment of mGluR 5 mediated disorders.

- 32. Use of the compound according to any one of claims 1 to 27, in the manufacture of a medicament for the treatment of mGluR 5 mediated disorders.
- 33. A method of treatment of mGluR 5 mediated disorders, comprising administrering to a mammal, including man in need of such treatment, a therapeutically effective amount of the compound according to any one of claims 1 to 27.
 - 34. The method according to claim 33, for use in treatment of neurological disorders.
 - 35. The method according to claim 33, for use in treatment of psychiatric disorders.
- 36. The method according to claim 33, for use in treatment of chronic and acute pain disorders.
 - 37. The method according to claim 33, for use in treatment of gastrointestinal disorders.
 - 38. A method for inhibiting activation of mGluR 5 receptors, comprising treating a cell containing said receptor with an effective amount of the compound according to claim 1.